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diagram of AdRSVpHyde was illustrated in Fig. 1. The sequence of AdRSVpHyde is set forth in Figure 10 (SEQ ID NO: 5 and SEQ ID NO: 6).—

Please replace the paragraph beginning on page 77, line 32 with the following rewritten paragraph:

13.

--Characterization of cDNA: Sequencing-p-Hyde cDNA was originally obtained as a λZAP Uni XR clone, and was further subcloned into pBluescript SK vector through in vivo excision protocol as described (Stratagen, La Jolla, California). This double-stranded cDNA was further subjected for Dye Terminator Cycle Sequencing (Perkin Elmer, Foster City, California) using ABI 377 automatic DNA sequencer Version 3.0. The open reading frame of p-Hyde cDNA was determined using the DNA Strider program (Pasteur Institute, Paris). --

In the Claims:

Please amend the following claims to read as follows:

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1. (Amended) An isolated nucleic acid molecule which encodes a mammalian p-Hyde protein, including variants, analogs and mutants thereof, said nucleic acid molecule set forth in SEQ ID No. 1.

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7. (Amended) The isolated nucleic acid molecule of claim 1, wherein the nucleic acid is DNA, c-DNA or RNA.

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- 10. (Amended) The isolated nucleic acid molecule of claim 1, wherein the nucleic acid is labeled with a detectable marker.
- 11. (Amended) The isolated nucleic acid molecule of claim 10, wherein the detectable marker is a radioactive, colorimetric, luminescent, fluorescent or gold label.
- 12. (Amended) An oligonucleotide of at least 15 nucleotides capable of specifically hybridizing with the molecule of claim 1.

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- 17. (Amended) An antisense molecule capable of specifically hybridizing with the isolated nucleic acid molecule of claim 1.
 - 18. (Amended) A vector comprising the isolated nucleic acid molecule of claim 1.

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19. (Amended) The vector of claim 18, further comprising an regulatory element linked to the nucleic acid molecule.

- 20. (Amended) The vector of claim 18, wherein the regulatory element comprises a bacterial, yeast, insect or mammalian promoter.
- 21. (Amended) The vector of claim 20, wherein the vector is a plasmid, cosmid, yeast artificial chromosome (YAC), BAC artificial chromosome, adenovirus, adeno-associated virus, retrovirus, P1 bacteriophage or eukaryotic viral DNA.

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- 23. (Amended) The adenovirus vector of claim 22, wherein the adenovirus vector comprises an adenovirus genome wherein the p-Hyde gene is inserted within a deletion in the E1 and E3 region of the genome.
- 54. (Amended) The isolated nucleic acid molecule of claim 1 having at least 75% complementary to the nucleic acid sequence of SEQ ID NO: 1.

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- 55. (Amended) The isolated nucleic acid molecule of claim 1 having at least 85% complementary to the nucleic acid sequence of SEQ ID NO: 1.
- 56. (Amended) The isolated nucleic acid molecule of claim 1 having at least 95% complementary to the nucleic acid sequence of SEQ ID NO: 1.
- 57. (Amended) The isolated nucleic acid molecule of claim 1 as set forth in SEQ ID NO: 1.

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- 59. (Amended) The isolated nucleic acid molecule of claim 53, wherein the nucleic acid is cDNA or genomic DNA.
- 60. (Amended) The isolated nucleic acid molecule of claim 1 encoding an amino acid sequence having the sequence as set forth in SEQ ID NO: 2.

REMARKS

Claims 1, 7, 10-27, 54-57, 59 and 60 are pending in the application. Claims 1, 7, 10-27, 54-57, 59 and 60 have been rejected. Claims 1, 7, 10-12, 17-21, 23, 54-57, 59 and 60 have been amended. The amendments to the claims, specification and abstract contain no new matter. Therefore, Applicants respectfully request entry of the Amendment.